## IN THE CLAIMS

- 1-22. (Canceled)
- (Currently Amended) The method of Claim [[19]]26 wherein the antibody is a humanized, human or chimeric antibody.
- 24. (Currently Amended) The method of Claim [[19]]26, wherein said pharmaceutically acceptable carrier is saline, buffered saline or glucose in saline.
- 25. (Currently Amended) The method of Claim [[19]]26, wherein said pharmaceutically acceptable carrier is selected from the group consisting of solid support, liposomes and micro spheres.
- 26. (Currently Amended) The method of Claim 19A method of inhibiting malignant cell migration in a host having a malignancy which is melanoma by administration of a migration-inhibiting effective amount of a composition containing an anti-CCL25 antibody in a pharmaceutically acceptable carrier, wherein said antibody is administered directly to tumor tissue.
- 27. (Currently Amended) The method of Claim [[19]]26, wherein said antibody is administered directly to tumor bed during an invasive procedure.
  - 28-29. (Canceled)
- 30. (New) The method of Claim 26, wherein said anti-CCL25 antibody is administered at a dose range of 0.01-1000 mg/kg/day.
- (New) The method of Claim 26, wherein said anti-CCL25 antibody is administered at a dose range of 0.1-100 mg/kg/day.
  - 32. (New) The method of Claim 25, wherein said solid support is a sponge or gauze.

- 33. (New) The method of Claim 23, wherein said anti-CCL25 antibody is a human antibody.
- 34. (New) The method of Claim 23, wherein said anti-CCL25 antibody is a humanized antibody.
- 35. (New) The method of Claim 23, wherein said anti-CCL25 antibody is a chimeric antibody.
- 36. (New) The method of Claim 24, wherein said pharmaceutically acceptable carrier is saline.
- 37. (New) The method of Claim 24, wherein said pharmaceutically acceptable carrier is buffered saline.
- 38. (New) The method of Claim 24, wherein said pharmaceutically acceptable carrier is glucose in saline.